Type L # Hits
Hits Search Text DBs Time Comme Pefi nts Stamp
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DBs Stamp Time Comme Defi niti AT; PGPUB; 2003/03/1 PGPUB; 2003/03/1 NAT; PGPUB; 2003/03/1 NAT; PGPUB; 2003/03/1 NAT; PGPUB; 2003/03/1
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1	2003/03/	USPAT; US-PGPUB; EPO; JPO; DERWENT	((shirley adj bret.in.) or (hora adj maninder.in.)) and (7 or 8)	Ν	L15	BRS	15
03/03/ 15:55	2003,	USPAT; US-PGPUB; EPO; JPO; DERWENT	hora adj maninder.in.	18	L14	BRS	14
2003/03/ 6 15:55	δ N	USPAT; US-PGPUB; EPO; JPO; DERWENT	shirley adj bret.in.	12	L13	BRS	13
2003/03/ 6 15:39	の N	USPAT; US-PGPUB; EPO; JPO; DERWENT	8 same 10 _.	0	L12	BRS	12
2003/03/ 6 15:39	و 0 0	USPAT; US-PGPUB; EPO; JPO; DERWENT	6 same 10	0	L11	BRS	11
2003/03/ 6 15:38	60	USPAT; US-PGPUB; EPO; JPO; DERWENT	(human adj insulin-like adj growth adj factor adj 1) or IGF-1	2186	L10	BRS	10
2003/03/	60	USPAT; US-PGPUB; EPO; JPO; DERWENT	(succinate same buffer) same ((human adj insulin-like adj growth adj factor adj ((pharmaceutical or therapeutic\$2) adj composition)) or IGF-1)	2	Ľ9	BRS	9
Time Stamp		DBs	Search Text	Hits	# #	Туре	

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(FILE 'HOME' ENTERED AT 16:01:17 ON 16 MAR 2003)

FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA' ENTERED AT

16:01:41 ON 16 MAR 2003

- L1 24950 S (PHARMACEUTIC? OR THERAPEUTIC?) (W) COMPOSITION
- L2 2100 S SUCCINATE (P) BUFFER
- L3 625 S (SUCCINIC ACID) (P) BUFFER
- L4 2617 S L2 OR L3
- L5 4 S L1 (P) L4
- L6 4 DUPLICATE REMOVE L5 (0 DUPLICATES REMOVED)
- L7 135 S COMPOSITION (P) L4
- L8 92 DUPLICATE REMOVE L7 (43 DUPLICATES REMOVED)
- L9 8 S L8 (P) MM
- L10 8 S L9 NOT L6
- L11 20744 S (HUMAN INSULIN-LIKE GROWTH FACTOR 1) OR (IGF-1)
- L12 0 S L11 (P) (L5 OR L9)

 $^{=&}gt; \log y$

FILE 'HOME' ENTERED AT 16:01:17 ON 16 MAR 2003 => file medline caplus biosis embase scisearch agricola COST IN U.S. DOLLARS SINCE FILE TOTAL **ENTRY** SESSION FULL ESTIMATED COST 0.21 0.21 FILE 'MEDLINE' ENTERED AT 16:01:41 ON 16 MAR 2003 FILE 'CAPLUS' ENTERED AT 16:01:41 ON 16 MAR 2003 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS) FILE 'BIOSIS' ENTERED AT 16:01:41 ON 16 MAR 2003 COPYRIGHT (C) 2003 BIOLOGICAL ABSTRACTS INC. (R) FILE 'EMBASE' ENTERED AT 16:01:41 ON 16 MAR 2003 COPYRIGHT (C) 2003 Elsevier Science B.V. All rights reserved.

FILE 'SCISEARCH' ENTERED AT 16:01:41 ON 16 MAR 2003 COPYRIGHT (C) 2003 Institute for Scientific Information (ISI) (R)

FILE 'AGRICOLA' ENTERED AT 16:01:41 ON 16 MAR 2003

=> s (pharmaceutic? or therapeutic?) (w) composition 24950 (PHARMACEUTIC? OR THERAPEUTIC?) (W) COMPOSITION

=> s succinate (p) buffer 2100 SUCCINATE (P) BUFFER

=> s (succinic acid) (p) buffer 625 (SUCCINIC ACID) (P) BUFFER

=> s 12 or 13 2617 L2 OR L3

=> s 11 (p) 144 L1 (P) L4

=> duplicate remove 15 PROCESSING COMPLETED FOR L5

4 DUPLICATE REMOVE L5 (0 DUPLICATES REMOVED)

=> d 16 1-4 ibib abs

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:256088 CAPLUS

DOCUMENT NUMBER:

136:299709

TITLE:

Tocol-based compositions containing amiodarone

INVENTOR(S): Lambert, Karel J.; Kessler, Dean R.; Nienstedt, Andrew M.; Hartgraves, Greg A.; Constantinides, Panayiotis P.

Sonus Pharmaceuticals, Inc., USA

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPL	ICATION NO.	DATE
WO 2002026324	A2 20020	0404 WO 2	001-US30320	20010927
WO 2002026324	A3 20020	0704		
W: AE, AG,	AL, AM, AT,	AU, AZ, BA, BB	, BG, BR, BY,	BZ, CA, CH, CN,
CO, CR,	CU, CZ, DE,	DK, DM, DZ, EC	, EE, ES, FI,	GB, GD, GE, GH,
GM, HR,	HU, ID, IL,	IN, IS, JP, KE	KG, KP, KR,	KZ, LC, LK, LR,
LS, LT,	LU, LV, MA,	MD, MG, MK, MN	, MW, MX, MZ,	NO, NZ, PH, PL,
PT, RO,	RU, SD, SE,	SG, SI, SK, SL	, TJ, TM, TR,	TT, TZ, UA, UG,
US, UZ,	VN, YU, ZA,	ZW, AM, AZ, BY	, KG, KZ, MD,	RU, TJ, TM

```
RW: GH, GM, KE, LS, MW, M7 SD, SL, SZ, TZ, UG, ZW, AT, BE CH, CY, DE, DK, ES, FI, FR, G GR, IE, IT, LU, MC, NL, PT, SI TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                      A5 20020408
                                        AU 2001-94826
                                                          20010927
    AU 2001094826
                                       US 2000-235865P P 20000927
PRIORITY APPLN. INFO.:
                                       WO 2001-US30320 W 20010927
      ***Pharmaceutical***
                              ***compns*** . comprising amiodarone or one of
    its prodrugs or analogs and one or more tocols are disclosed. An emulsion
    contained amiodarone 0.6, d,l-.alpha.-tocopherol 1.0, tocopherol
    polyethylene glycol ***succinate*** 1.0, Poloxamer P-407 0.5, PEG-400
             ***buffer*** q.s. 50 mL. Stability of the formulations was
    g, and
     studied.
    ANSWER 2 OF 4 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                    1996:121156 CAPLUS
DOCUMENT NUMBER:
                        124:156044
                       Pharmaceutical compositions containing hGH.
TITLE:
INVENTOR(S):
                       Samaritani, Fabrizio
                       Applied Research Systems, Neth.
PATENT ASSIGNEE(S):
                        PCT Int. Appl., 25 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
    PATENT NO.
                   KIND DATE
                                        APPLICATION NO. DATE
                                         ______
     WO 9535116
                    A1 19951228
                                        WO 1994-IT86
                                                         19940617
        W: JP, US
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
                                        EP 1994-920573
                                                         19940617
     EP 804223 A1 19971105
                          19990922
                     B1
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE
                                    JP 1994-501903 19940617
    JP 10504531 T2 19980506
                                         AT 1994-920573
    AT 184798
                      F.
                           19991015
                                                          19940617
    ES 2139081
                     T3
                                        ES 1994-920573
                           20000201
                                                          19940617
                                        US 1996-750684 19961217
    US 5898030
                     A 19990427
                                      EP 1994-920573
PRIORITY APPLN. INFO.:
                                                          19940617
                                       WO 1994-IT86
                                                          19940617
     Pharmaceutical compns. contq. hGH were stabilized by saccharose. The
     formulation is particularly suitable for stabilizing a lyophilizate of
    recombinant hGH.
    ANSWER 3 OF 4 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER:
                    1995:532058 CAPLUS .
DOCUMENT NUMBER:
                        122:274053
TITLE:
                        Process and apparatus for manufacturing of a
                        pharmaceutical composition containing prednisolone
                        sodium succinate, suitable for parenteral dosing
INVENTOR(S):
                        Mago Karacsony, Erzsebet; Ambrus, Gabor; Balogh,
                        Tibor; Danitz, Bela; Toldy, Lajos; Makk, Nandor;
                        Tegdes, Aniko; Kovacs, Klara Maria; Bidlo, Gaborne; et
PATENT ASSIGNEE(S):
                        Gyogyszerkutato Intezet, Hung.
SOURCE:
                        Hung. Teljes, 14 pp.
                        CODEN: HUXXBU
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        Hungarian
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                   KIND DATE
                                        APPLICATION NO. DATE
                                         -----
     -----
    HU 66012 A2 19940829
HU 212306 B 19960528
                                        HU 1992-4081
                                                         19921222
PRIORITY APPLN. INFO.:
                                       HU 1992-4081
                                                         19921222
    The process involves mixing prednisolone hemisuccinate and NaOH, sterile
    filtering of the resultant prednisolone sodium succinate soln., filling it
    into ampuls, lyophilizing it, and closing the ampuls under an inert gas
     atm. Thus, powd. prednisolone hemisuccinate with a particle size
```

.ltoreq.200 .mu.m is dispersed in an aq. soln. contg. (9.5.+-.0.2):(0.5.+-

.0.2) wt.:wt. Na2HPO4 and NaH2O4 as buffer substances. The dispersion is cooled to 5-15.degree., prefer by to 5-10.degree. Then 80-5 preferably 85-95%, of the stoichiometrically necessary 0.3-1.0% wt.:vol. NaOH soln. is added in portions during intensive stirring of the reaction medium and stirring is continued until the complete dissoln. of prednisolone hemisuccinate. A stainless steel reactor for carrying out the process is also claimed. In contrast to former processes this process gives only trace amts. of hydrolysis products at most.

L6 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1990:62635 CAPLUS
DOCUMENT NUMBER: 112:62635
TITLE: Stabilized injection solutions containing nonlyophilized gamma-interferons
INVENTOR(S): Hwang-Felgner, Jiin Yu; Jones, Richard E.; Maher, James F.

PATENT ASSIGNEE(S): Genentech, Inc., USA SOURCE: PCT Int. Appl., 15 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

```
PATENT NO.
                    KIND DATE
                                          APPLICATION NO. DATE
     WO 8904177 A1 19890518
                                           WO 1988-US3883 19881101
         W: AU, DK, FI, HU, JP, KR, NO
         RW: AT, BE, CH, DE, FR, GB, IT, LU, NL, SE
     IL 88233
                    A1 19930818 IL 1988-88233
                                                             19881030
                      A1 19890601
                                           AU 1988-27245
     AU 8827245
                                                             19881101
     AU 621327
                      B2 19920312
    EP 386106 A1 19900912
EP 386106 B1 19940302
                                           EP 1988-910211
                                                             19881101
        R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE

    JP 03500882
    T2 19910228

    JP 2732877
    B2 19980330

                                      JP 1988-509401
                                                             19881101
                     E 19940315
                                          AT 1988-910211 19881101
     AT 102048
                     A 19900725
A5 19910502
     ZA 8808249
                                          ZA 1988-8249
                                                            19881103
     DD 289470
                                          DD 1988-321429 19881103
     CA 1335176 A1 19950411
US 5151265 A 19920929
                                      CA 1988-582102 19881103
US 1990-514392 19900425
PRIORITY APPLN. INFO.:
                                        US 1987-116434
                                                            19871103
                                        EP 1988-910211 19881101
WO 1988-US3883 19881101
```

AB A liq. ***pharmaceutical*** ***compn*** . comprises an effective amt. of nonlyophilized .gamma.-interferon. The compn. further includes a ***buffer*** capable of maintaining the pH within 4-6, polyhydric sugar alcs. as stabilizer, and a nonionic detergent. The relative shelf-life for the liq. contg. 2 mg/mL .gamma.-interferon, mannitol, and ***succinate*** ***buffer*** was 10 days as compared to 1 day for the lyophilized formulation.

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L2

L3

L4 L5 (FILE 'HOME' ENTERED AT 16:01:17 ON 16 MAR 2003)

FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA' ENTERED AT 16:01:41 ON 16 MAR 2003

24950 S (PHARMACEUTIC? OR THERAPEUTIC?) (W) COMPOSITION

2100 S SUCCINATE (P) BUFFER

625 S (SUCCINIC ACID) (P) BUFFER

2617 S L2 OR L3

4 S L1 (P) L4

=> s composition (p) 14 L7 135 COMPOSITION (P) L4

=> duplicate remove 17
DUPLICATE PREFERENCE IS 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH'

4 DUPLICATE REMOVE L5 (0 DUPLICATES REMOVED)

KEEP DUPLICATES FROM MORE THAN ONE_LE? Y/(N):n PROCESSING COMPLETED FOR L7

92 DUPLICATE REMOVE L7 (43 DUPLICATES REMOVED)

=> s 18 (p) mM

PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'L49 (P) MM' PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'L55 (P) MM'

8 L8 (P) MM

=> s 19 not 16

8 L9 NOT L6 L10

=> d l10 1-8 ibib abs

L10 ANSWER 1 OF 8 MEDLINE

88307898 ACCESSION NUMBER: MEDLINE

DOCUMENT NUMBER: 88307898 PubMed ID: 2457334

Separation of cell organelles in density gradients based on TITLE:

their permeability characteristics.

Gasser K W; DiDomenico J; Hopfer U AUTHOR:

Department of Physiology and Biophysics, Case Western CORPORATE SOURCE:

Reserve University, Cleveland, Ohio 44106.

CONTRACT NUMBER: AM 25170 (NIADDK)

> DK 27651 (NIDDK) HL 07415 (NHLBI)

ANALYTICAL BIOCHEMISTRY, (1988 May 15) 171 (1) 41-6. SOURCE:

Journal code: 0370535. ISSN: 0003-2697.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE) DOCUMENT TYPE:

LANGUAGE: English

Priority Journals FILE SEGMENT:

ENTRY MONTH: 198809

ENTRY DATE: Entered STN: 19900308

Last Updated on STN: 19970203 Entered Medline: 19880915

The buoyant density of intracellular organelles is dependent in part on AB the nature of the ***buffer*** ***composition*** of the density gradient and the permeability characteristics of the organelle membrane to the constituents of this ***buffer*** . Therefore, knowledge of the transport properties of different organelles allows the design of density gradients useful for their purification. We have used this approach to significantly decrease mitochondrial contamination of pancreatic zymogen granules in a one-step purification procedure on a 40% Percoll density gradient. These gradients, prepared with isoosmotic sucrose, yield a narrow band of zymogen granules and mitochondria. However, by substitution of sucrose with salts to which mitochondria but not zymogen granules are permeable, the densities of mitochondria are altered to give a significant separation. For example, the incorporation of 100 ***MM***

succinate in the Percoll gradient can produce a 70% reduction in mitochondrial contamination. The increased ionic strength has an additional beneficial effect on zymogen granule yield by 5-10%. The recognition and utilization of transport pathways in organelle membranes is the principal feature of this technique and should prove to be widely

applicable to other isolation procedures.

L10 ANSWER 2 OF 8 MEDLINE

ACCESSION NUMBER: 83238332 MEDLINE

DOCUMENT NUMBER: 83238332 PubMed ID: 6305947

TITLE: The regulation of extramitochondrial steady state free Ca2+

concentration by rat insulinoma mitochondria.

AUTHOR: Prentki M; Janjic D; Wollheim C B

JOURNAL OF BIOLOGICAL CHEMISTRY, (1983 Jun 25) 258 (12) SOURCE:

7597-602.

Journal code: 2985121R. ISSN: 0021-9258.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198308

ENTRY DATE: Entered STN: 19900319 Last Updated or STN: 19970203 Entered Medlin 19830811

For the study of Ca2+ handling by mitochondria of an insulin secretory tissue, a method for the isolation of functionally intact insulinoma mitochondria is described. The mitochondria had a respiratory control ratio of 6.3 +/- 0.3 with ***succinate*** as a substrate. The regulation of extramitochondrial [Ca2+]o concentration by suspensions of insulinoma mitochondria was studied using Ca2+-selective minielectrodes. The mitochondria were found to maintain an ambient free Ca2+ concentration of about 0.3 and 0.9 microM in the absence or presence of Mg2+ (1 ***mM***), respectively. The addition of Na+ resulted in a dose-dependent (half-maximal 4 ***mM*** Na+) increase in steady state [Ca2+]o. Na+ accelerated the ruthenium red-induced Ca2+ efflux, suggesting the existence of a Ca2+/2Na+ antiporter, as described in mitochondria of excitable tissues. Experiments were performed to study the effects of various agents on the steady state extramitochondrial free Ca2+. cAMP, 3-isobutyl-1-methylxanthine, and NADH were found to have no effect,

whereas phosphoenolpyruvate induced a net Ca2+ efflux, the kinetic of which suggests deleterious effects on mitochondrial functions. A small decrease in pH (0.1 unit) of the incubation ***buffer*** resulted in an increase of the extramitochondrial Ca2+ steady state that was reversible upon restoration of the pH to its initial value. In conclusion, insulinoma mitochondria were able to maintain an extramitochondrial [Ca2+]o steady state in the submicromolar range that was markedly influenced by the ionic ***composition*** of the incubation medium. Thus, mitochondria may play a role in the regulation of cellular calcium homeostasis and insulin release.

L10 ANSWER 3 OF 8 MEDLINE

ACCESSION NUMBER: 83230736 MEDLINE

DOCUMENT NUMBER: PubMed ID: 6860312 83230736

Photosynthetic electron transport in thylakoid preparations TITLE:

from two marine red algae (Rhodophyta).

AUTHOR: Stewart A C; Larkum A W

SOURCE: BIOCHEMICAL JOURNAL, (1983 Feb 15) 210 (2) 583-9.

Journal code: 2984726R. ISSN: 0264-6021.

PUB. COUNTRY: ENGLAND: United Kingdom

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198307

ENTRY DATE: Entered STN: 19900319

> Last Updated on STN: 19900319 Entered Medline: 19830729

Thylakoid membrane preparations active in photosynthetic electron AB transport have been obtained from two marine red algae, Griffithsia monilis and Anotrichium tenue. High concentrations (0.5-1.0 M) of salts such as phosphate, citrate, ***succinate*** and tartrate stabilized functional binding of phycobilisomes to the membrane and also stabilized Photosystem II-catalysed electron-transport activity. High concentrations (1.0 M) of chloride and nitrate, or 30 ***mM*** -Tricine/NaOH

(pH 7.2) in the absence of salts, detached phycobilisomes ***buffer*** and inhibited electron transport through Photosystem II. The O2-evolving system was identified as the electron-transport chain component that was inhibited under these conditions. Washing membranes with containing 1.0-1.5 M-sorbitol and 5-50 ***mM*** concentrations of various salts removed the outer part of the phycobilisome but retained 30-70% of the allophycocyanin 'core' of the phycobilisome. These preparations were 30-70% active in O2 evolution compared with unwashed membranes. In the sensitivity of their O2-evolving apparatus to the

composition of the medium in vitro, the red algae resembled blue-green algae and differed from other eukaryotic algae and higher plants. It is suggested that an environment of structured water may be essential for the functional integrity of Photosystem II in biliprotein-containing algae.

L10 ANSWER 4 OF 8

ACCESSION NUMBER: 76260253 MEDLINE

DOCUMENT NUMBER: 76260253 PubMed ID: 783158

TITLE: Effect of cations and anions on the steady state kinetics

of energy-dependent Ca2+ transport in rat liver

mitochondria.

MEDLINE

AUTHOR: Hutson S M; Pfeiffer D R; Lardy H A

SOURCE: JOURNAL OF BIO ICAL CHEMISTRY, (1976 Sep 10) (17)

5251-8.

Journal code: 2985121R. ISSN: 0021-9258.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 197611

ENTRY DATE: Entered STN: 19900313

Last Updated on STN: 19970203 Entered Medline: 19761101

The divalent cation ionophore A23187 has been used to investigate the kinetics of energy-dependent Ca2+ uptake by rat liver mitochondria under steady state conditions. During A23187-induced cyclic Ca2+ flux, the free Ca2+ concentration is adjusted using [ethylenebis(oxyethylenenitrilo)]tetr aacetic acid (EGTA) ***buffers*** . The rate of Ca2+ transport, which is inferred from the rate of ***succinate*** oxidation, is a function of the free Ca2+ concentration in the medium. The kinetics are sigmoidal with the free Ca2+ concentration at half-maximal respiratory stimulation (K0.5) equal to 3.1 +/- 0.4 muM at 25 degrees. The maximal Ca2+-stimulated respiratory rate (Vmax) is a function of the ionic ***composition*** of the medium. Magnesium and Mg2+ plus phosphate produced a parallel stimulation of the maximal respiration rate whether activated by Ca2+ uptake or by the uncoupler carbonyl cyanide-ptrifluoromethoxyphenylhydrazone (FCCP). In the absence of A23187, Ca:O rations of 4.0 were obtained under most experimental conditions. Magnesium is a potent competitive-like inhibitor, increasing the K0.5 for Ca2+ to 30.0 muM at 2.0 ***mM*** MgCl2. Magnesium dramatically decreases the apparent affinity for Ca2+ but does not appear to alter the kinetic mechanism. In contrast, the alkali metal cations are weak inhibitors, at most doubling the K0.5 for Ca2+; however, they antagonized Mg2+ inhibition with an order of effectiveness Li+ greater than or equal to Na+ greater than K+ greater than Rb+ =Cs+. Phosphate and acetate increased the Vmax slightly without altering the K0.5 for Ca2+. Phosphate did not influence the inhibitory effects of Mg2+ or Mg2+ plus K+. This study suggests that during steady state conditions, the maximal rate of Ca2+ accumulation is primarily electron transport-limited. The results are also discussed in terms of a possible physiological role for Mg2+ and K+ in the intracellular regulation of energy-dependent mitochondrial Ca2+ transport in liver.

L10 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1994:79106 CAPLUS

DOCUMENT NUMBER: 120:79106

TITLE: Manufacture of transparent moldings with hard surface INVENTOR(S): Uenishi, Michiharu; Nagai, Shoichi; Takei, Masatoshi;

Kobayashi, Yukio; Akaqi, Juji

PATENT ASSIGNEE(S):

Mitsubishi Rayon Co, Japan Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

SOURCE:

Patent Japanese

LANGUAGE:
FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 05228431 A2 19930907 JP 1992-32341 19920219
PRIORITY APPLN. INFO.: JP 1992-32341 19920219

Title moldings, useful for building, automotive, and optical applications (no data), are manufd. by irradiating moldings bearing on surface a layer of polymers derived from monomers contg. .gtoreq.2 (meth)acryloyl groups and other monomers with UV light of wavelengths .ltoreq.300 nm, treating the irradiated moldings with alkali to generate .gtoreq.0.02 .mu.mol acidic groups/cm2 on the surface, coating the surface with hydrolytic polycondensation products of XaSi(OY)4-a (X = epoxy-contg. functional group; Y = hydrocarbyl; a = 1-3) and/or SiRlbR2c(OR3)d [R1, R2 = (ether or ester linkage-contg.) hydrocarbyl; R3 = H, hydrocarbyl; b, c = 0-3; d = 4 - b - c = 1-4; b + c = 1-3], and contacting the polycondensation products with a high-temp. fluid at .gtoreq.500.degree. Thus, a 2 ***mm*** -thick PMMA sheet was dip-coated with a soln. contg. dipentaerythritol

hexaacrylate 10, equimolar **succinic*** ***acid***
-trimethylolethane-acrylic accondensation product 20,
tetrahydrofurfuryl acrylate 5, Darocur 1173 1.2, isopropanol 34, and
toluene 20% and irradiated with UV (365 nm, 840 mJ/cm2) at 35.degree. to
form a 3.5 .mu.m-thick film with pencil hardness 5H, which was further
UV-irradiated (254 nm, 1300 mJ/cm2) and immersed in 20% aq. NaOH to
generate 0.05 .mu.mol acidic groups/cm2. The sheet was dip-coated with a
compn . of .gamma.-glycidoxypropyltrimethoxysilane 100.4,
isopropanol 278.3, tetraethoxysilane 40.0, 0.2 N AcOH/NaOAc (
buffer , pH 5) 22.9, and Mg perchlorate 2.0 parts, held at
100.degree. for 3 h, then brought into contact with a natural gas flame of
900.degree. 20 times for .apprx.0.2 s each time. The cured coat showed
good Taber abrasion resistance, cross-cut adhesion 100/100 initially and
100/100 after 20-h immersion in H2O at 80.degree., and smooth crack-free
surface before and after the hot water immersion.

L10 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1989:639540 CAPLUS

DOCUMENT NUMBER: 111:239540

TITLE: Liposomes containing hydrophilic drugs and a process

for manufacture them

INVENTOR(S): Profitt, Richard Thomas; Adler-Moore, Jill; Chiang,

Su-Ming

PATENT ASSIGNEE(S): Vestar, Inc., USA

SOURCE: Eur. Pat. Appl., 13 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	TENT NO.	KIND	DATE	APPLICATION NO. DATE
•		A1 B1		EP 1988-310278 19881101
	R: AT, BE,	CH, DE	, ES, FR,	GB, GR, IT, LI, LU, NL, SE
AU	8824161	A1	19890518	AU 1988-24161 19881024
AU	598958	B2	19900705	·
AT	66598	E	19910915	AT 1988-310278 19881101
ES	2029330	Т3	19920801	ES 1988-310278 19881101
KR	9707187	B1	19970507	KR 1988-14547 19881105
NO	8804989	Α	19890516	NO 1988-4989 19881109
NO	178484	В	19960102	
ИО	178484	C	19960410	
JP	01160915	A2	19890623	JP 1988-284828 19881110
JP	2958774	B2	19991006	
CA	1339008	A1	19970325	CA 1988-582730 19881110
DK	8806293	Α	19890513	DK 1988-6293 19881111
US	5965156	Α	19991012	US 1995-469251 19950606
PRIORIT	Y APPLN. INFO.	. :		US 1987-119518 A 19871112
				EP 1988-310278 A 19881101
				US 1990-600154 A1 19901019
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A novel liposome ***compn*** . and a method for solubilizing AB amphiphilic drugs in a small amt. of org. solvent for use in improved liposomes are described. A phosphatidylglycerol is acidified and the amphiphilic drugs suspended in an org. solvent are added to solubilize the drugs. Distearoylphosphatidylglycerol Na soln. dissolved in CHCl3-MeOH mixt. (1:1) was acidified with HCl and then mixed with amphotericin B (I) soln. dissolved in the same solvent. Hydrogenated egg phosphatidylcholine soln. and cholesterol soln. dissolved in the same solvent were then mixed with the mixt. The pH was adjusted to 4.5 by addn. of 2.5 N NaOH. molar ratio of I, distearoylphosphatidylglycerol, hydrogenated egg phosphatidylcholine, and cholesterol in the soln. was 0.4, 0.4, 2.0, and 1.0 resp. The lipid soln. was spray-dried to give a powder, which was hydrated with 9% lactose-contg. 10 ***mM*** ***succinate***

buffer (pH 5.62) and sonicated to give liposomes. Mice were i.v. inoculated with Candida albicans and 3 days post-infection, mice were treated with a single dose of either free I or liposomal I. There was no dose level of free I which produced any survivors at 29 days post-infection; however, all animals treated with 10 or 15 mg/kg of liposomal I were still alive 42 days post-infection.

L10 ANSWER 7 OF 8 CAPLUS COPYRICHT 2003 ACS ACCESSION NUMBER: 1989:5807 CAPLUS

DOCUMENT NUMBER: 111:180738

TITLE: Sustained-release pharmaceuticals containing a soluble metoprolol salt and a dihydropyridine in a gel-forming

matrix

INVENTOR(S): Ragnarsson, Gert Anders; Silfverstrand, Kajsa

Margareta; Sjoegren, John Albert

PATENT ASSIGNEE(S): Aktiebolag Haessle, Swed. SOURCE: Eur. Pat. Appl., 11 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA'	rent no.	KIND	DATE		APPLICATION NO.	DATE
			19890412		EP 1988-850319	19880922
EP		B1				~=
		•		GB, G	R, IT, LI, LU, NL	•
AU	8822374	A1	19890413		AU 1988-22374	19880916
AU	615211	B2	19910926			
AT	84412	E	19930115		AT 1988-850319	19880922
ES	2053815	Т3	19940801		ES 1988-850319	19880922
NO	8804269	Α	19890410		NO 1988-4269	19880927
NO	177375	В	19950529			
NO	177375	С	19950906			
US	4942040	Α	19900717		US 1988-250945	19880929
IL	87922	A1	19930708		IL 1988-87922	19881005
DK	8805586	A	19890409		DK 1988-5586	19881006
FI	8804636	Α	19890409		FI 1988-4636	19881007
FI	92903	В	19941014			
FI	92903	С	19950125			
JР	01128917	A2	19890522		JP 1988-252209	19881007
CA	1312286	A1	19930105		CA 1988-579566	19881007
	1032490	A	19890426		CN 1988-109129	19881008
=	1029935	В	19951011			
	Y APPLN. INFO).:		SE	1987-3881	19871008
111201111					1988-850319	19880922
					1700 030317	17000722

AB A controlled-release pharmaceutical for once daily administration contains metoprolol and a poorly water-sol. Ca channel blocking agent of the dihydropyridine type; metoprolol is included in the form of small beads contg. as the main sol. component a salt of metoprolol coated with a water-insol. polymeric membrane. The dihydropyridine is dispersed in a nonionic solubilizer. Both, the dispersed dihydropyridine and the metoprolol-contg. beads are incorporated in a matrix forming a swelling agent when in contact with water. A mixt. contg. felodipine, Polyoxyl-40 hydrogenated castor oil (solubilizer), Polyvydon-K90, hydroxypropyl Me cellulose (swelling agent), Al silicate, lactose, and microcryst. cellulose was granulated with EtOH and dried. Metoprolol

succinate was sprayed onto cores of SiO2 to form beads (0.5 diam.) and the beads were coated by spraying with a soln. contg. Et cellulose, hydroxypropyl Me cellulose in CH2Cl2/iso-PrOH; the beads and granules were mixed, a lubricant was added and the . was pressed into tablets. Tablets contg. 10 mg felodipine and 95 mg metoprolol ***succinate*** each were prepd. from a mixt. contg. felodipine 10, Polyoxyl-40 25, Polyvydon-K90 24, hydroxypropyl Mg cellulose 230, Al silicate 94, lactose 56, microcryst. cellulose 6, ***succinate*** 95, SiO2 24, Et cellulose 32, and addl. hydroxypropyl Me cellulose 8 g. The dissoln. rate of felodipine in phosphate ***buffer*** contg. 1% Na dodecyl sulfate was 14, 64, 88, and 98% after 2, 8, 12, and 20 h, resp.; the dissoln. rater of metorpolol ***succinate*** was 5, 39, 65, and 95% after 2, 8, 12, and 20 h, resp. Suitable polymers for coating the beads are Eudragit RL, Eudragit RS, alone or in combination; Et cellulose in combination with hydroxypropyl Me cellulose or hydroxypropyl cellulose is preferred.

L10 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1987:118245 CAPLUS

DOCUMENT NUMBER: 106:118245

TITLE: Freshness of fish and shellfish. II. Modified simple

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colorimetry of dehydrogenase activity in shellfish using tet colium chloride
                         Tsunoda, Kojun; Inoue, Noriko; Aoyama, Mitsuo; Hasebe,
AUTHOR (S):
                         Akihisa
                         Suginami Ward Inst. Public Health, Tokyo, 168, Japan
CORPORATE SOURCE:
SOURCE:
                         Shokuhin Eiseigaku Zasshi (1986), 27(5), 487-91
                         CODEN: SKEZAP; ISSN: 0015-6426
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         Japanese
     A simple colorimetry procedure for the detn. of dehydrogenase [9035-82-9]
     activity in shellfish was based on redn. of triphenyltetrazolium chloride
     (TTC) to triphenylformazan (TF). The dehydrogenase activity in shellfish
     muscle is low. When pH 7.4 phosphate
                                             ***buffer*** was added to the
     TTC reagent, dehydrogenase activity in shellfish muscle was increased
     markedly. The best ***compn*** . of TTC reagent was 0.2% TTC-0.1% Na
       ***succinate*** -2.84% Na2HPO4-2.34% NaCl. A sample of shellfish muscle
     was sliced about 1.5 ***mm*** thick, soaked in TTC reagent, incubated
     at 37.degree. for 30 min, cooled, extd. with EtOH, filtered, and measured
     at 284 nm. It was also possible to det. dehydrogenase activity in oyster
     gills by using this TTC reagent.
=> d his
     (FILE 'HOME' ENTERED AT 16:01:17 ON 16 MAR 2003)
     FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA' ENTERED AT
     16:01:41 ON 16 MAR 2003
L1
          24950 S (PHARMACEUTIC? OR THERAPEUTIC?) (W) COMPOSITION
L2
           2100 S SUCCINATE (P) BUFFER
            625 S (SUCCINIC ACID) (P) BUFFER
L3
L4
           2617 S L2 OR L3
L5
              4 S L1 (P) L4
L6
              4 DUPLICATE REMOVE L5 (0 DUPLICATES REMOVED)
L7
            135 S COMPOSITION (P) L4
L8
             92 DUPLICATE REMOVE L7 (43 DUPLICATES REMOVED)
L9
              8 S L8 (P) MM
              8 S L9 NOT L6
=> s (human insulin-like growth factor 1) or (iqf-1)
   4 FILES SEARCHED...
         20744 (HUMAN INSULIN-LIKE GROWTH FACTOR 1) OR (IGF-1)
=> s l11 (p) (15 or 19)
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L73 (P) '
PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L76 (P) '
             0 L11 (P) (L5 OR L9)
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     16:01:41 ON 16 MAR 2003
L1
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L2
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L3
            625 S (SUCCINIC ACID) (P) BUFFER
           2617 S L2 OR L3
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L5
              4 S L1 (P) L4
L6
              4 DUPLICATE REMOVE L5 (0 DUPLICATES REMOVED)
L7
            135 S COMPOSITION (P) L4
             92 DUPLICATE REMOVE L7 (43 DUPLICATES REMOVED)
L8
L9
              8 S L8 (P) MM
              8 S L9 NOT L6
L10
L11
          20744 S (HUMAN INSULIN-LIKE GROWTH FACTOR 1) OR (IGF-1)
              0 S L11 (P) (L5 OR L9)
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=> log yCOST IN U.S. DOLLARS